

AD	

GRANT NO: DAMD17-94-J-4143

TITLE: Sodium MRI of Rat Breast Tumors

PRINCIPAL INVESTIGATOR(S): Hee Kwon Song

CONTRACTING ORGANIZATION: University of Pennsylvania

Philadelphia, Pennsylvania

19104-3246

REPORT DATE: September 1995

TYPE OF REPORT: Annual

PREPARED FOR:

Commander

U.S. Army Medical Research and Materiel Command

Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for public release;

distribution unlimited

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

19951211 095

## REPORT DOCUMENTATION PAGE

Form Approved
OMB No. 0704-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204. Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503.

Davis nightvay, suite 1204, Armigion, VA 222	02-4502, and to the office of Management and	ounger, reper nor necessarion respect (or or o	,
1. AGENCY USE ONLY (Leave bla	ank) 2. REPORT DATE 199	5 3. REPORTIVE 1AND PAIS	94¥8731/95)
4. TITLE AND SUBTITLE	t Droogt Tumora	5. FUN	IDING NUMBERS
Sodium MRI of Ra	it Breast Tumors	D.F	MD17-94-J-4143
6. AUTHOR(S)			
Hee Kwon Song			
T DEDECTRAINE ODCANIZATION	MANUE(C) AND ADDRESS(ES)	I O DED	FORMING ORGANIZATION
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) University of Pennsylvania			ORT NUMBER
Philadelphia, Pennsylvania 19104-3246			
111111111111111111111111111111111111111			
9. SPONSORING/MONITORING A	GENCY NAME(S) AND ADDRESS(ES	) 10. SPC AG	DNSORING/MONITORING ENCY REPORT NUMBER
	-1 Deserved and Mat	arrial Command	
	al Research and Mat aryland 21702-5012		
Fort Detrick, Ma	iryrand 21702-3012		
11. SUPPLEMENTARY NOTES	er i o plane solito, como de conserva e emandado do desenção y o entreja de sendamente administração do habitado em conserva de conserva d		entre de la companya
12a. DISTRIBUTION / AVAILABILITY	STATEMENT	12b. D	ISTRIBUTION CODE
Approved for pub	olic release; distr	bution unlimited	
Approved for pas	110 1010000, 01001.		
	47		
13. ABSTRACT (Maximum 200 wor	rds) to use sodium MR imag	: 41; 4. 4: <i>66</i>	1
tumor types in the r	at. Specifically, we prop	ong techniques to difference to determine and continuous	ompare the sodium
concentration and the	T1 and T2 relaxation para	meters for two rat breast	tumor lines. Since
sodium images have i	ntrinsically low signal-to-no	oise ratios (SNR), the first	part of this research
involves implementing	g various methods to increat higher field MR magnet, b	se the SNR of sodium ima	ges. Three methods
	a surface coil, and implem		
	struction. This part of the		
	successfully, and the MR so		ned to collect data to
	the backprojection technique multiexponential decays		imple apostrogeony
	rmed on three rats implanted		
visible in the data, al	though there were large va	riations in the values of	the decay constants,
	usceptibility effects. This e		
	ters from the actual imaging	experiments since spin ec	hoes will be used for
imaging.			
14. SUBJECT TERMS	dootion wasanst-	hronet imagina	15. NUMBER OF PAGES
Sodium imaging, projection reconstruction, breast imaging		, preast imaging	16. PRICE CODE
breast cancer			
17. SECURITY CLASSIFICATION OF REPORT	18. SECURITY CLASSIFICATION OF THIS PAGE	19. SECURITY CLASSIFICATION OF ABSTRACT	20. LIMITATION OF ABSTRACT
Ünclassified	OF THIS PAGE Unclassified	OF ABSTRACT Unclassified	Unlimited

#### **GENERAL INSTRUCTIONS FOR COMPLETING SF 298**

The Report Documentation Page (RDP) is used in announcing and cataloging reports. It is important that this information be consistent with the rest of the report, particularly the cover and title page. Instructions for filling in each block of the form follow. It is important to stay within the lines to meet optical scanning requirements.

- Block 1. Agency Use Only (Leave blank).
- **Block 2.** Report Date. Full publication date including day, month, and year, if available (e.g. 1 Jan 88). Must cite at least the year.
- Block 3. Type of Report and Dates Covered. State whether report is interim, final, etc. If applicable, enter inclusive report dates (e.g. 10 Jun 87 30 Jun 88).
- Block 4. <u>Title and Subtitle</u>. A title is taken from the part of the report that provides the most meaningful and complete information. When a report is prepared in more than one volume, repeat the primary title, add volume number, and include subtitle for the specific volume. On classified documents enter the title classification in parentheses.
- Block 5. Funding Numbers. To include contract and grant numbers; may include program element number(s), project number(s), task number(s), and work unit number(s). Use the following labels:

C - Contract PR - Project
G - Grant TA - Task
PE - Program WU - Work Unit
Element Accession No.

- **Block 6.** <u>Author(s)</u>. Name(s) of person(s) responsible for writing the report, performing the research, or credited with the content of the report. If editor or compiler, this should follow the name(s).
- **Block 7.** <u>Performing Organization Name(s) and Address(es).</u> Self-explanatory.
- Block 8. <u>Performing Organization Report</u>
  <u>Number</u>. Enter the unique alphanumeric report number(s) assigned by the organization performing the report.
- **Block 9.** Sponsoring/Monitoring Agency Name(s) and Address(es). Self-explanatory.
- Block 10. Sponsoring/Monitoring Agency Report Number. (If known)
- Block 11. Supplementary Notes. Enter information not included elsewhere such as: Prepared in cooperation with...; Trans. of...; To be published in.... When a report is revised, include a statement whether the new report supersedes or supplements the older report.

Block 12a. <u>Distribution/Availability Statement</u>. Denotes public availability or limitations. Cite any availability to the public. Enter additional limitations or special markings in all capitals (e.g. NOFORN, REL, ITAR).

DOD - See DoDD 5230.24, "Distribution Statements on Technical Documents."

DOE - See authorities.

NASA - See Handbook NHB 2200.2.

NTIS - Leave blank.

Block 12b. Distribution Code.

DOD - Leave blank.

DOE - Enter DOE distribution categories from the Standard Distribution for Unclassified Scientific and Technical Reports.

NASA - Leave blank. NTIS - Leave blank.

- Block 13. Abstract. Include a brief (Maximum 200 words) factual summary of the most significant information contained in the report.
- **Block 14.** <u>Subject Terms</u>. Keywords or phrases identifying major subjects in the report.
- **Block 15.** <u>Number of Pages</u>. Enter the total number of pages.
- Block 16. <u>Price Code</u>. Enter appropriate price code (NTIS only).
- Blocks 17. 19. Security Classifications. Self-explanatory. Enter U.S. Security Classification in accordance with U.S. Security Regulations (i.e., UNCLASSIFIED). If form contains classified information, stamp classification on the top and bottom of the page.
- Block 20. <u>Limitation of Abstract</u>. This block must be completed to assign a limitation to the abstract. Enter either UL (unlimited) or SAR (same as report). An entry in this block is necessary if the abstract is to be limited. If blank, the abstract is assumed to be unlimited.

#### FOREWORD

Opinions, interpretations, conclusions and recommendations are those of the author and are not necessarily endorsed by the US Army.

Where copyrighted material is quoted, permission has been obtained to use such material.

Where material from documents designated for limited distribution is quoted, permission has been obtained to use the material.

Citations of commercial organizations and trade names in this report do not constitute an official Department of Army endorsement or approval of the products or services of these organizations.

In conducting research using animals, the investigator(s) adhered to the "Guide for the Care and Use of Laboratory Animals, prepared by the Committee on Care and Use of Laboratory Animals of the Institute of Laboratory Resources, National Research Council (NIH Publication No. 86-23, Revised 1985).

For the protection of human subjects, the investigator(s) adhered to policies of applicable Federal Law 45 CFR 46.

In conducting research utilizing recombinant DNA technology, the investigator(s) adhered to current guidelines promulgated by the National Institutes of Health.

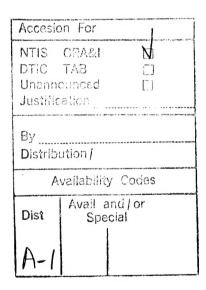
In the conduct of research utilizing recombinant DNA, the investigator(s) adhered to the NIH Guidelines for Research Involving Recombinant DNA Molecules.

In the conduct of research involving hazardous organisms, the investigator(s) adhered to the CDC-NIH Guide for Biosafety in Microbiological and Biomedical Laboratories.

He Kwow Som/ Sept. 29, 1945
PI - Signature Date

# **Table of Contents**

<u>Page</u>	<u>Item</u>
1	Front Cover
2	SF 298 Report Documentation Page
3	Foreword
4	Table of Contents
5	Introduction
6	Methods and Developments
7	Results and Conclusions
8	References
10	Appendix A



#### Introduction

Reliable diagnosis of breast tumors is still in its developmental stages. Currently, mammography is widely used as a breast screening test, but is often unsuccessful in detecting and diagnosing tumors. It is difficult to detect tumors in women with dense breasts. Also, cancers that are located high in the breast and close to the chest wall are hard to detect with mammography. In addition, even if a lesion is discovered, it is difficult to determine whether it is malignant or benign since there is a considerable overlap in the tumors' shapes and densities. Biopsies are often necessary to determine the malignancy of detected tumors.

Magnetic resonance imaging (MRI) is becoming more popular as a diagnostic tool in breast cancer research. Although in the earlier years MRI had little to offer in the diagnosis of breast tumors over the less expensive mammogram, it is becoming more popular and promising as newer techniques are developed. The use of gadolinium (Gd) contrast agents has shown that malignant breast tumors consistently enhance, while benign lesions, such as scars and fat necrosis, do not. [1] More recently, "dynamic" contrast-enhanced MRI has been investigated. [2-8] In these studies, the investigators study not only the enhancement of the tumors, but also the rate of enhancement as contrast agents are injected into the patients. Studies show that cancerous tissue generally enhances faster than benign tumors. Some authors indicate, however, that these differences are not yet statistically significant and that other factors should also be considered in the final diagnosis. [7] Although studies with Gd-contrast agents have demonstrated that contrast-enhanced proton MRI has excellent sensitivity that exceeds that of mammography, the specificity still varies considerably. [9] Even with the use of these agents, it is still difficult to differentiate carcinomas from fibroadenomas; both tumor types often enhance by similar amounts. [10] Although MRI is more sensitive than mammography, used alone, it is not fully adequate for determining the malignancy of tumors.

The goal of this research is to investigate a different method to detect and characterize different types of breast tumors. It has been known over the years that sodium concentration is significantly higher in tumor cells than in normal cells. [11-17] X-ray microanalysis, NMR spectroscopy, and MRI experiments have demonstrated the increase in sodium (23Na) content in neoplastic tissue. Because various cellular activities, including mitosis and oncogenesis, seem to depend heavily on cellular ionic concentrations and content, especially that of sodium, 23Na MR imaging is potentially a more sensitive imaging technique than the conventional proton imaging method for both detection and differentiation of tumors. Sodium studies have been done in the past in both human and animal models, but no one has yet looked specifically into sodium MR imaging of different breast tumors and its potential to differentiate tumors without the need for more invasive techniques. The purpose of this research is to determine the sodium characteristics, including the concentration and T1 and T2 relaxation parameters, for two different types of rat breast tumors, 13762A and Ac33.

Sodium images have intrinsically low signal-to-noise ratios (SNR). Therefore, to acquire reliable quantitative data, we will combine three methods to increase the SNR of our images: use a higher field magnet; use a surface coil rather than a body coil during the receive mode; and implement a projection reconstruction technique to reduce the echo time. All of these techniques will aid in improving the SNR and in detecting all of the T1 and T2 components of the sodium signal.

### **Methods and Developments**

The first step in improving the SNR of sodium images is by using a higher field MRI machine. A 4.7 Tesla animal imaging system has been installed in our laboratory. Previously, most of the animal studies have been done on the 1.9 Tesla animal imager. We can expect the signal-to-noise ratio to increase by a factor of about 2.5 with the new system, since SNR increases linearly with increasing field. We also have a six-inch inner diameter gradient set for the high field magnet. The set up is complete and has been proven to give high quality images in animal studies.

The second step was to build an RF coil that optimizes the SNR of our images. An RF coil pair has been constructed for this purpose. A linearly polarized birdcage coil for transmitting houses a smaller surface coil used to receive the signal. Since a smaller receive coil will pick up less noise than a larger one while being more sensitive to the region of interest, we can expect a significant increase in SNR. In order to decouple the two coils, they are oriented such that the rf field that each of them produces is orthogonal to the other. The surface coil is further decoupled from the transmit coil actively using a diode as described in literature. [18] The isolation between the two coils during RF transmission is 56 dB, and during receive (when decoupling is only passive) is 26 dB. Good images have been obtained in test studies.

The third way to increase the image quality is by reducing the echo time of our pulse sequence. Since the sodium signal decays exponentially after the spins have been excited, signal strength is strongest just after the RF pulse. Also, a short echo time is necessary in order to collect the fast component of T2 of sodium. To accomplish this, a pulse program for the MRI scanner implementing the backprojection algorithm was written and tested. The program consists of an initial 90 degree hard pulse followed by data collection. There is no phase encoding step as in conventional Fourier imaging. The data collection starts immediately following the RF pulse and is sampled with a constant interval throughout the readout period. During this period the gradients are in transition, and it is therefore necessary to interpolate the data points for accurate reconstruction. The program also collects data from subsequent spin echoes.

In order to reconstruct the collected data, a 3-dimensional filtered backprojection reconstruction algorithm was written in Pascal on the PC. The algorithm was tested successfully on both simulated data and real data from phantom imaging experiments. A simple linear interpolation was performed for data taken from the imaging experiments.

As a quick test for biexponential decay from sodium signals from the tumors, sodium spectroscopy was performed on the tumors in vivo. Ac33 rat mammary tumors were transplanted bilaterally into three rats. After two to three weeks the rats were anesthetized and spectroscopy was performed on the tumors. For two of the rats a single surface coil was used for spectroscopy, while for the third rat the new RF coil pair described above was used. Table 1 summarizes the results obtained from the experiments.

#### **Results and Conclusions**

The main goal of this period was to improve the quality (specifically, increase the SNR) of sodium images. RF coils optimized for sodium imaging of breast tumors were built, and filtered backprojection pulse program for the MRI scanner was implemented. Further work will involve modifying the sodium body coil to be able to obtain proton images as well. A well-known double tuning method using "tank circuits" can be used to accomplish this. [19]

From the Table 1, one can see that the standard deviations of the T2 values obtained from the spectroscopy experiments are large. This is probably due to local field inhomogeneities caused by susceptibility difference between air and tissue. Susceptibility effects will not be a major issue in the actual imaging experiments since spin echoes will be used to obtain the data for echoes two and above. The images from the first set of data points collected from the FID will also not be affected by susceptibility effects since the SNR depends primarily on the first few points of the data, which is collected before significant signal decay occurs.

The focus of the first year of the grant was on the implementation and construction of various tools necessary for successful sodium imaging experiments. Nearly all of the hardware and software needed to image rat breast tumors were built and written during this period.

#### References

- [1] Harms, S.E. and D.P. Flaming, MR imaging of the breast, J. Magn. Reson. Imag. 3, 277-283 (1993).
- [2] Kaiser, W.A. and E. Zeitler, MR imaging of the breast: Fast imaging sequences with and without Gd-DTPA, Radiology 170, 681-686 (1989).
- [3] Stack, J.P., O.M. Redmond, M.B. Codd, P.A. Dervan, and J.T. Ennis, Breast disease: Tissue characterization with Gd-DTPA enhancement profiles, Radiology 174, 491-494 (1990).
- [4] Turner, D.A., J.Z. Wang, S.G. Economou, M. Cobleigh, K.J. Bloom, T.R. Witt, and E. Staren, Functional images from dynamic, contrast-enhanced, 3DFT MR images for the detection of breast cancer, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 116 (1993).
- [5] Heβ, T., M.V. Knopp, G. Brix, U. Hoffmann, H. Junkermann, H.-J. Zabel, and G. van Kaick, Pharmacokinetic mapping of breast lesion by dynamic Gd-DTPA enhanced MRI, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 117 (1993).
- [6] Weisskoff, R.M., C.A. Hulka, B. Smith, K. McCarthy, D.A. Hall, G.J. Whitman, D.B. Kopans, and T.J. Brady, Dynamic NMR imaging of the breast using echo planar imaging, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 119 (1993).
- [7] Schnall, M.D., S. Orel, and L. Muenz, Analysis of time intensity curves for enhancing breast lesions, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 120 (1993).
- [8] Kelcz, F., G.E. Santyr, S.J. Mongin, and E.J. Fairbanks, Reducing false positive gadolinium-enhanced breast MRI results through parameter analysis of the enhancement profile, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 121 (1993).
- [9] Heywang-Kobrunner, S.H., MRI of breast disease, Proc. of Twelfth Annual Meeting of Society of Magnetic Resonance in Medicine, 113 (1993).
- [10] Heywang, S.H., A. Wolf, E. Pruss, T. Hilbertz, W. Eiermann, and W. Permanetter, MR imaging of the breast with Gd-DTPA: Use and limitations, Radiology 171, 95-103 (1989).
- [11] Goldsmith, M. and R. Damadian, NMR in cancer. VII. Sodium-23 magnetic resonance of normal and cancerous tissues, Physiol. Chem. Phys. 7, 263-269 (1975).
- [12] Cameron, I.L., N.K.R. Smith, T.B. Pool, and R.L. Sparks, Intracellular concentration of sodium and other elements as related to mitogenesis and oncogenesis in vivo, Canc. Res. 40, 1493-1500 (1980).
- [13] Shen, S.S, S.T. Hamamoto, H.A. Bern, and R.A. Steinhardt, Alteration of sodium transport in mouse mammary epithelium associated with neoplastic transformation,

- Canc. Res. 38, 1356-1361 (1978).
- [14] Zs.-Nagy, I, G. Lustyik, G. Lukacs, V. Zs.-Nagy, and G. Balazs, Correlation of malignancy with the intracellular Na+:K+ ratio in human thyroid tumors, Canc. Res. 43,5395-5402 (1983).
- [15] Gupta, R.K., 23Na NMR spectroscopy of intact cells and tissues. In R.K. Gupta (ed): NMR Spectroscopy of Cells and Organisms, vol 2. Boca Raton: CRC Press, Inc., pp. 1-32 (1987).
- [16] Hilal, S.K., J.B. Ra, C.H. Oh, I.K. Mun, S.G. Einstein, and P. Roschmann, Sodium Imaging. In D.D. Stark and W.G. Bradley (eds): Magnetic Resonance Imaging. St. Louis: CV Mosby, pp. 715-731 (1988).
- [17] Summers, R.M., P.M. Joseph, and H.L. Kundel, Sodium nuclear magnetic resonance imaging of neuroblastoma in the nude mouse, Invest. Radiol. 26, 233-241 (1991).
- [18] Edelstein, W.A., C.J. Hardy, and O.M. Mueller, Electronic decoupling of surface-coil receivers for NMR imaging and spectroscopy, J. Magn. Reson. 67, 156-161 (1986).
- [19] Schnall, M.D., V.H Subramanian, J.S. Leigh, J. Magn. Reson. 67, 129-134 (1986).

# Appendix A

**Table 1** Slow and fast components of sodium T2 values and their standard deviations in implanted rat breast tumors using MR spectroscopy.

	Average (ms)	Standard Deviation (ms)
T2 fast	2.4	1.13
T2 slow	8.5	5.60